

**What is claimed is:**

1.

A method of screening animals to determine those more likely to produce larger litters comprising:  
obtaining a sample of genetic material from said animal; and  
assaying for the presence of a genotype in said animal which is associated with increased litter size, said genotype characterized by the following:

a) a polymorphism in the PRKAG3 gene.

2.

The method of claim 1 wherein said polymorphism results in an amino acid change from valine to isoleucine at amino acid number 199 of the PRKAG3 gene or its equivalent as determined by a BLAST comparison of SEQ ID NO:2.

3.

The method of claim 1 wherein said polymorphism is a transition of a guanine to an adenine at nucleotide position 595 or its equivalent.

4.

The method of claim 1 wherein said genotype is a BsaHI polymorphism.

5.

The method of claim 1 wherein said step of assaying is selected from the group consisting of:  
restriction fragment length polymorphism (RFLP) analysis, minisequencing, MALD-TOF, SINE, heteroduplex analysis, single strand conformational polymorphism (SSCP), denaturing gradient gel electrophoresis (DGGE) and temperature gradient gel electrophoresis (TGGE).

6.

The method of claim 1 wherein said animal is a pig.

7.

The method of claim 1 further comprising the step of amplifying the amount of PRKAG3 gene or a portion thereof which contains said polymorphism.

8.

The method of claim 7 wherein said amplification includes the steps of:  
selecting a forward and a reverse sequence primer capable of  
amplifying a region of the PRKAG3 gene which contains a  
polymorphic BsaHI site.

9.

The method of claim 8 wherein said forward and reverse  
primers are selected from and based upon primer RNF and  
primer RNR.

10.

A method of screening animals to determine those more  
likely to exhibit improved meat quality traits comprising:  
obtaining a biological sample of material from said animal;  
and  
assaying for the presence of a genotype in said animal which  
is associated with improved meat quality traits said  
genotype characterized by the following:

- a) a polymorphism in the PRKAG3 gene, said  
polymorphism resulting in and characterized by an  
amino acid of valine at position 199 and arginine  
at position 200, or an isoleucine at position 199  
when an arginine is at position 200 or its  
equivalent as determined by a BLAST comparison of  
SEQ ID NO:2.

11.

The method of claim 10 wherein said polymorphism is a  
transition of a guanine to an adenine at nucleotide position  
595 or its equivalent.

12.

The method of claim 10 wherein said step of assaying  
comprises a short interspersed element polymorphism test.

13.

The method of claim 12 wherein said assay comprises the  
step of amplifying the PRKAG3 gene using primers selected  
from and based upon primer RP1F and primer PN52R2.

14.

The method of claim 10 further comprising the step of amplifying the amount of PRKAG3 gene or a portion thereof which contains said polymorphism.

15.

The method of claim 14 wherein said amplification includes the steps of:  
selecting a forward and a reverse sequence primer capable of amplifying a region of the PRKAG3 gene which contains a polymorphic BsaHI site.

16.

The method of claim 14 wherein said forward and reverse primers are selected from and based upon Primer RNF and primer RNR.

17.

A method of screening animals to determine those more likely to exhibit improved meat quality traits comprising:  
obtaining a biological sample of material from said animal;  
and  
assaying for the presence of a genotype in said animal which is associated with improved meat quality traits said genotype characterized by the following:

- a) a polymorphism in the PRKAG3 gene, said polymorphism resulting in and characterized by an amino acid change of asparagine to threonine at amino acid position 30 or its equivalent as determined by a BLAST comparison of SEQ ID NO:1.

18.

The method of claim 17 wherein said polymorphism is a transition of an adenine to cytosine at nucleotide position 89 or its equivalent as determined by a BLAST comparison of SEQ ID NO:1.

19.

The method of claim 17 wherein said genotype is a Styl polymorphism.

20.

The method of claim 17 wherein said step of assaying is selected from the group consisting of:

restriction fragment length polymorphism (RFLP) analysis, minisequencing, MALD-TOF, SINE, heteroduplex analysis, single strand conformational polymorphism (SSCP), denaturing gradient gel electrophoresis (DGGE) and temperature gradient gel electrophoresis (TGGE).

21.

The method of claim 20 wherein said animal is a pig.

22.

The method of claim 20 further comprising the step of amplifying the amount of PRKAG3 gene or a portion thereof which contains said polymorphism.

23.

The method of claim 22 wherein said amplification includes the steps of:

selecting a forward and a reverse sequence primer capable of amplifying a region of the PRKAG3 gene which contains a polymorphic StyI site.

24.

The method of claim 23 wherein said forward and reverse primers are selected from and based upon Primer RF1 and primer RN52R2.

25.

A method of screening animals to determine those more likely to exhibit improved meat quality traits comprising: obtaining a biological sample of material from said animal; and assaying for the presence of a genotype in said animal which is associated with improved meat quality traits said genotype characterized by the following:

- a) a polymorphism in the PRKAG3 gene, said polymorphism resulting in and characterized by an amino acid change of glycine to serine at amino acid position 52 or its equivalent as determined by a BLAST comparison of SEQ ID NO:1.

26.

The method of claim 25 wherein said polymorphism is a transition of a guanine to an adenine at nucleotide position 154 or its equivalent as determined by a BLAST comparison of SEQ ID NO:1.

27.

The method of claim 25 wherein said genotype is a HphI polymorphism.

28.

The method of claim 25 wherein said step of assaying is selected from the group consisting of:

restriction fragment length polymorphism (RFLP) analysis, minisequencing, MALD-TOF, SINE, heteroduplex analysis, single strand conformational polymorphism (SSCP), denaturing gradient gel electrophoresis (DGGE) and temperature gradient gel electrophoresis (TGGE).

29.

The method of claim 25 wherein said animal is a pig.

30.

The method of claim 28 further comprising the step of amplifying the amount of PRKAG3 gene or a portion thereof which contains said polymorphism.

31.

The method of claim 30 wherein said amplification includes the steps of:  
selecting a forward and a reverse sequence primer capable of amplifying a region of the PRKAG3 gene which contains a polymorphic HphI site.

32.

The method of claim 30 wherein said forward and reverse primers are selected from and based upon Primer RF1 and primer RN52R2

33.

A nucleotide sequence which encodes upon expression an PRKAG3 protein, further comprising a serine at position 52.

34.

The nucleotide sequence of claim 33 comprising SEQ ID NO:5.

35.

A PRKAG3 protein according to claim 33.

36.

The protein of claim 35 comprising SEQ ID NO:6.

37.

A nucleotide sequence which encodes upon expression an PRKAG3 protein, said protein comprising a isoleucine at position 199 and an arginine at position 200 or the equivalent thereof, of said protein.

38.

A PRKAG3 protein according to claim 37.

39.

A nucleotide sequence which encodes upon expression an PRKAG3 protein, said protein comprising an isoleucine at position 199, a threonine at position 30 a glycine at position 52 and an arginine position 200 or the equivalent thereof, of said protein.

40.

A PRKAG3 protein according to claim 39.

41.

A nucleotide sequence which encodes upon expression an PRKAG3 protein, said protein comprising a valine at position 199 and an arginine at position 200 or the equivalent thereof, of said protein.

42.

A PRKAG3 protein according to claim 41.

43.

A nucleotide sequence which encodes upon expression an PRKAG3 protein, said protein comprising an isoleucine or valine at position 199 and an arginine at position 200 or the equivalent thereof, of said protein.

44.

A PRKAG3 protein according to claim 43.

45.

A method of screening animals to determine those more likely to have favorable meat quality traits comprising: obtaining a sample of genetic material from said animal; and assaying for the presence of a genotype in said animal which is associated with favorable meat quality traits, said genotype characterized by the following:  
a threonine at amino acid position 30, a glycine at amino acid position 52 and an isoleucine at amino acid position 199.

46.

A method of screening animals to determine those more likely to have favorable meat quality traits comprising: obtaining a sample of genetic material from said animal; and assaying for the presence of a genotype in said animal which is associated with favorable meat quality traits, said genotype characterized by the following:  
a isoleucine at position 199 and an arginine at position 200.

47.

A method for identifying a genetic marker for meat quality and/or litter size in animals comprising the steps of:  
determining the number of offspring produced by each female animal or the meat quality of said animal;  
determining the polymorphism in the PRKAG3 or equivalent gene of each animal; said polymorphism comprising the polymorphism of claim 1, 7, 17, or 11 or their equivalents and  
associating the number of offspring produced by each female animal or meat quality with said polymorphism thereby identifying a polymorphism for animal meat quality or litter size.

48.

The method of claim 47 further comprising the step of selecting animals for breeding which are predicted to have favorable meat quality or litter size by said marker.

Table 1. Demographic characteristics of the study population	
Age (years)	18-24
	25-34
	35-44
	45-54
Sex	Male
	Female
	Male
	Female
Marital status	Married
	Single
	Divorced
	Widowed
Education level	High school or less
	Some college
	Bachelor's degree
	Postgraduate
Occupation	Professional
	Managerial
	Service
	Unemployed
Income (USD/month)	<1000
	1000-1999
	2000-2999
	≥3000
Health status	Good
	Fair
	Poor
	Very poor
Smoking status	Never
	Former
	Current
	Unknown
Alcohol consumption	Never
	Former
	Current
	Unknown
Exercise frequency	Never
	1-2 times/week
	3-4 times/week
	≥5 times/week

50.

51.

52.

53.

a) a polymorphism in the PRKAG3 gene, said polymorphism being one other than the RN<sup>-</sup> mutation at amino acid 200.



54.

A method of screening animals to determine those more likely to have favorable meat quality traits comprising: obtaining a sample of genetic material from said animal; and assaying for the presence of a genotype in said animal which is associated with favorable meat quality, said genotype characterized by a combination of at least two polymorphisms in the PRKAG3.

55.

A method of screening animals to determine those more likely to have increased value for litter size and/or meat quality traits comprising: obtaining a sample of genetic material from said animal; and assaying for the presence of a genotype in said animal which is associated with favorable litter size and/or meat quality, said genotype characterized by a combination of at least two polymorphisms in the PRKAG3.

56.

A method of screening animals to determine those more likely to have favorable meat quality traits comprising: obtaining a sample of genetic material from said animal; and assaying for the presence of a genotype in said animal which is associated with favorable meat quality traits, said genotype characterized by the following:  
a threonine at amino acid position 30, a serine at amino acid position 52 and a valine at amino acid position 199.

57.

A method of screening animals to determine those more likely to exhibit improved meat quality traits and or larger litter size comprising: obtaining a biological sample of material from said animal; and  
assaying for the presence of a genotype in said animal which is associated with said traits said genotype characterized by the following:

- a) a short interspersed element polymorphism in the PRKAG3 gene.

58.

The method of claim 57 wherein said assay comprises the step of amplifying the PRKAG3 gene using primers selected from and based upon primer RP1F and primer PN52R2.

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